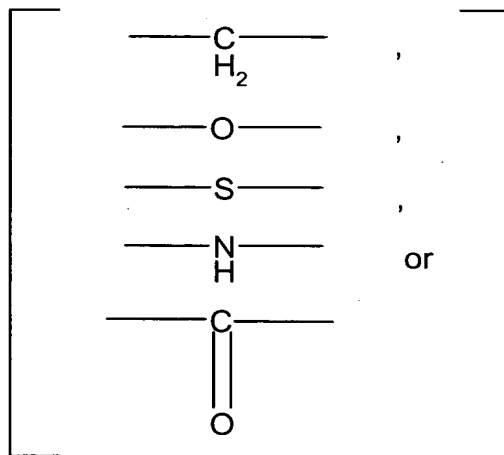


Amendments to the Claims

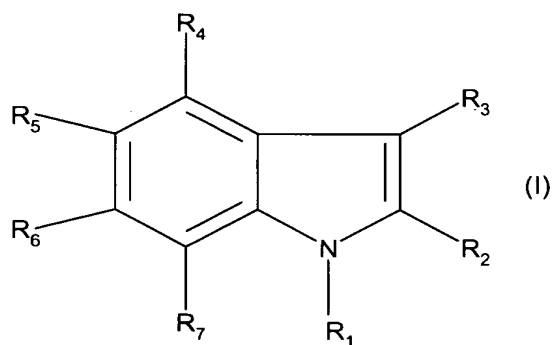
This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended)



An indole compound represented by the formula (I), or a pharmaceutically acceptable salt, solvate, or prodrug derivative thereof;



wherein;

R_1 is selected from groups (a), (b), and (c) wherein;

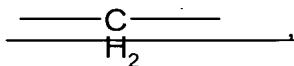
(a) is C7-C20 alkyl, C7-C20 haloalkyl, C7-C20 alkenyl, C7-C20 alkynyl, carbocyclic radical, or heterocyclic radical, or

(b) is a member of (a) substituted with one or more independently selected non-interfering substituents; or

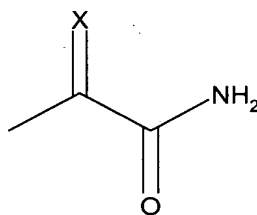
(c) is the group $-(L_1)-R_{11}$; where, $-(L_1)-$ is a divalent linking group of 1 to 8 atoms and where R_{11} is a group selected from C7-C20 alkyl, C7-C20 haloalkyl, C₁-C₄alkylaryl, or aryl;

R_2 is hydrogen, or a C1-C45 alkyl;

R_3 is $-(L_3)-Z$, where $-(L_3)-$ is a ~~a bond~~ bond, or



and Z is selected from a group represented by the formulae,



wherein, X is oxygen;

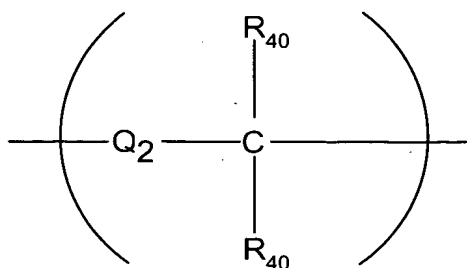
R_4 is the group, $-(L_h)-$ (hydroxyfunctional amide); wherein $-(L_h)-$, is an hydroxyfunctional amide linker having an hydroxyfunctional amide linker length of 1 to 8;

R_5 is the group, $-(L_a)-$ (acidic group); wherein $-(L_a)-$, is an acid linker having an acid linker length of 1 to 8;

R_6 and R_7 are each independently selected from hydrogen, C₁-C₆ alkyl, C₂-C₆alkenyl, and C₂-C₆ alkynyl.

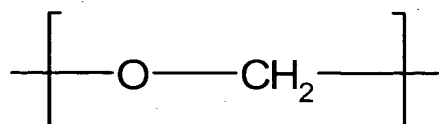
2. (Previously Amended) The compound of claim 1 wherein R_2 is hydrogen, or C₁-C₄ alkyl.

3. (Previously Amended) The compound of Claim 1 wherein the hydroxyfunctional amide linker group, $-(L_h)-$, for R_4 is selected from a group represented by the formula;

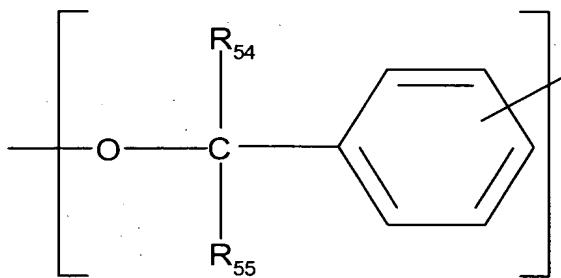


where Q₂ is -O-, and each R₄₀ is independently selected from hydrogen, and C₁-C₈ alkyl.

4. (Original) The compound of Claim 1 wherein the hydroxyfunctional amide linker group, -(Lh)-, for R₄ is a divalent group selected from,

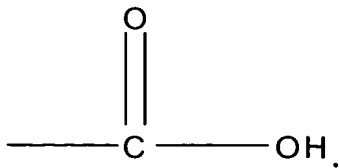


5. (Previously Amended) The compound of Claim 1 wherein the acid linker, -(L_a)-, for R₅ is the group represented by the formula;



wherein R₅₄, and R₅₅, are each independently hydrogen, or C₁-C₈ alkyl.

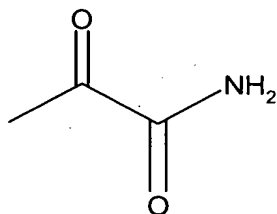
6. (Previously Amended) The compound of claim 1 wherein R₅ is the group, -(L_a)-(acidic group) and wherein the (acidic group) is the group:



7. (Canceled)

8. (Canceled)

9. (Original) The compound of claim 1 wherein for R₃, Z is the group represented by the formula;



and the linking group $-(L_3)-$ is a bond.

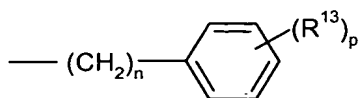
10. (Canceled)

11. (Canceled)

12. (Canceled)

13. (Canceled)

14. (Previously Amended) The compound of claim 1 wherein the linking group - (L₁₁)- of R₁₁ is a bond and R₁₁ is -(CH₂)_m-R¹² wherein m is an integer from 1 to 6, and R¹² is a group represented by the formula:



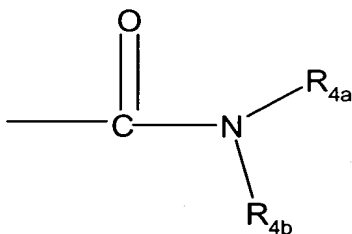
wherein a, and p are independently an integer from 0 to 2.

15. (Canceled)

16. (Canceled)

17. (Canceled)

18. (Previously Amended) The compound of claim 1 wherein R₄ is the group, -(L_C)-(hydroxyfunctional amide group) and wherein the (hydroxyfunctional amide group) is:



and R^{4a} is the ~~group OH~~group OH; and

wherein R^{4b} is independently selected from the group consisting of H, and (C₁-C₆)alkyl.

19. (Canceled)

20. (Canceled)

21. (Canceled)

22. (Original) A pharmaceutical formulation comprising a indole compound as claimed in claim 1 together with a pharmaceutically acceptable carrier or diluent therefor.

23. (Previously Amended) A method of inhibiting sPLA₂ mediated release of fatty acid comprising contacting sPLA₂ with a therapeutically effective amount of indole compound as claimed in claim 1.

24. (Currently Amended) A method of treating ~~a mammal~~ rheumatoid arthritis in a mammal; ~~to alleviate the pathological effects of Inflammatory Diseases~~; wherein the method comprises administering to said mammal a therapeutically effective amount of an indole compound as claimed in Claim 1.

25. (Canceled)

26. (Canceled)

27. (Canceled)